

Pregnancy Rates Among Adolescents Silicosis Screening in Surface Coal

Miners — Pennsylvania, 1996–1997 616 Hepatitis B Vaccination Coverage Among Asian and Pacific Islander Children — United States, 1998

619 Delayed Supply of Influenza Vaccine and Adjunct ACIP Influenza Vaccine Recommendations for the 2000–01 Influenza Season

622 Summary of the Joint Statement on Thimerosal in Vaccines

# National and State-Specific Pregnancy Rates Among Adolescents — United States, 1995–1997

Each year in the United States, 800,000–900,000 adolescents aged ≤19 years become pregnant. Adolescent pregnancy and childbearing have been associated with adverse health and social consequences for young women and their children. This report presents estimated national numbers of pregnancies and national and state-specific pregnancy rates for adolescents aged ≤19 years from 1995\* to 1997. The findings indicate a decline in national and state-specific adolescent pregnancy rates during 1995–1997, and a continuing downward trend beginning in the early 1990s (1,2,4).

Number of pregnancies was estimated as the sum of live births, legally induced abortions, and estimated fetal losses (i.e., spontaneous abortions and stillbirths) among females aged ≤19 years. Live birth data were reported by the mother's state of residence. Because abortion data by residence were not available for all states, abortions were reported by state of occurrence.¹ Complete age-specific abortion information was not available for nine reporting areas in 1995 (including the District of Columbia [DC]), eight areas in 1996 (including DC), and six states in 1997. To calculate national adolescent pregnancy rates, estimates of abortions among adolescents were calculated for states with missing data (1). Estimates of fetal losses were based on sample survey data of women aged 15–44 years from the 1988 and 1995 National Surveys of Family Growth (NSFG) (3). A national estimate of fetal losses for all females aged 15–19 years was derived from NSFG data and used to create annual estimates of fetal losses based on the number of live births and legally induced abortions in a given year (CDC, unpublished

<sup>\*</sup>National and state-specific adolescent pregnancy rates for 1995 were previously reported (1,2). National rates for 1995 are reported here because fetal loss estimates were not included in the earlier definition of pregnancy (1) and because of a change in the population denominator data supplied by the Bureau of the Census used in calculating rates; state-specific data for 1995 are reported again because of the change in the population denominator data. Adolescent pregnancy rates previously published by CDC (2) should not be used together with those reported here in time series analyses because of these changes in methods. Adolescent pregnancy rates in other sources (3) may not be comparable to data in this report because of different calculation methodologies.

For 48 reporting areas in 1995–1996 and 49 in 1997, the number and characteristics of persons who had legal induced abortions were provided by state health departments and the health departments of New York City and the District of Columbia. For four areas in 1995–1996 and three in 1997, the number of abortions were provided from hospitals and other medical facilities.

data, 1998). Denominators (estimates of the adolescent female population by state, age, and race) for abortion and fetal loss rates were obtained from postcensal population estimates. Published birth rates were added to abortion and fetal loss rates and were based on earlier, slightly different population estimates (5).

Rates were calculated as the number of pregnancies per 1000 females aged 15–17, 18–19, or 15–19 years. Because most pregnancies, births, and abortions (97% of live births and 94% of legally induced abortions) among females aged <15 years occurred among 13–14-year-olds (CDC, unpublished data, 2000; 6), this age group was used as the denominator for calculating rates for females aged <15 years. Legally induced abortions for which mother's age or race was unknown were included in categories based on the distribution of mothers with known age or race.

Although abortion totals were available for all states, age-specific data adequate to calculate pregnancy\*\* rates were available from 42 states and DC for 1995, 44 states and DC in 1996, and 45 states and DC in 1997. Because adequate age and Hispanic ethnicity data for females who had abortions were available for 24 states in 1995 (7), 23 states in 1996, and 26 states in 1997, pregnancy rates by ethnicity were not included; some states with missing Hispanic ethnicity data had large Hispanic populations.

From 1995 to 1997, among females aged 15–19 years, the national number of pregnancies declined by 3.1% and the national pregnancy rate declined by 7.8%, from 98.3 per 1000 in 1995 to 90.7 in 1997 (Table 1). During 1995–1997, the pregnancy rate declined by 11.3% among females aged <15 years, by 10.7% among females aged 15–17 years, and by 5.8% among females aged 18–19 years. For each year, the pregnancy rate for 18–19-year-olds was approximately 2.5 times that of 15–17-year-olds, and the rate for females aged <15 years was approximately one ninth that of 15–17-year-olds.

TABLE 1. Estimated number of pregnancies\* and rates' among adolescents, by age and year — United States, 1995–1997

	Estim	ated no. o	of pregnan	cies		Pregnanc	y rate	
Year	<15	15-17	18-19	15-19	<15	15-17	18-19	15-19
1995	26,600	342,100	525,000	867,100	7.2	63.9	151.4	98.3
1996	25,400	332,500	526,700	859,200	6.8	60.5	147.8	94.8
1997	23,700	321,300	518,800	840,000	6.4	57.1	142.7	90.7
% Change from	n							
1995 to 19976	-11.1%	-6.1%	-1.2%	-3.1%	-11.3%	-10.7%	-5.8%	-7.8%

<sup>\*</sup> Rounded to the nearest 100.

Available on the World-Wide Web at http://www.census.gov/population/estimates/state/ 5age9890.txt. Accessed July 2000.

Birth rates for females aged <15 years were calculated using 13-14-year-olds as the denominator.

<sup>\*\*</sup> Pregnancy rates were excluded if they were based on <20 pregnancies or <1000 adolescents in a particular category, or if >15% of the pregnancies were in women of unknown age or

Per 1000 adolescent females in the appropriate age group (per 13-14-year-olds for <15 years age group). For states that did not report abortion data by age (nine in 1995, eight in 1996, and six in 1997), numbers of abortions were estimated.</p>

<sup>&</sup>lt;sup>5</sup> Percent changes were computed on the basis of unrounded numbers and rates.

State-specific pregnancy rates per 1000 among 15–19-year-olds ranged¹¹ from 56.3 (North Dakota) to 117.1 (Nevada) in 1995; from 53.9 (North Dakota) to 114.1 (Texas) in 1996; and from 48.2 (North Dakota) to 127.8 (Delaware) in 1997 (Table 2). In each year, the rate for each reporting area was highest for females aged 18–19 years and lowest for females aged <15 years. From 1995 to 1997, the pregnancy rate for 15–19-year-olds decreased in 40 of the 43 reporting areas for which age-specific data were available. Statistically significant declines occurred in 34 states and ranged from 1.9% (Ohio) to 19.8% (Maryland); no state showed a significant increase. During 1995–1997, significant declines in the pregnancy rate occurred among females aged <15 years in 20 of 41 reporting areas with available data, among 15–17-year-olds in 35 of 42 reporting areas, and among 18–19-year-olds in 27 of 42 reporting areas.

Pregnancy rates for 15–19-year-olds were, in every state except one, higher for blacks than for whites among the 30 states with available data for both groups (Table 3). Significant declines in the pregnancy rate occurred among whites in 29 of the 35 states for which adequate data for whites were available, and in 17 of 28 states for which adequate data for blacks were available. No significant increases in pregnancy rates were found for adolescents of either race in states with available data.

Among females aged 15–19 years, the national birth rate decreased from 56.8 in 1995 to 52.3 in 1997 (*5*), with declines occurring in most reporting areas. The national number of abortions declined 2.7% from 1995 to 1997, and the national abortion rate decreased 7.4%, from 26.6 per 1000 in 1995 to 24.6 in 1997. During this period, the abortion rate decreased 3.9% among females aged <15 years (from 2.8 to 2.7), 10.1% among females aged 15–17 years (from 18.2 to 16.3), and 5.4% among females aged 18–19 years (from 39.6 to 37.5). From 1995 to 1997, the abortion rate for 15–19-year-olds decreased in 32 of the 43 reporting areas for which age-specific data were available. In 25 of the 31 areas where both birth and abortion rates decreased, the percent decrease in abortion rates exceeded the decline in birth rates.

Reported by: Behavioral Epidemiology and Demographic Research Br and Statistics and Computer Resources Br, Div of Reproductive Health, National Center for Chronic Disease Prevention and Health Promotion, CDC.

Editorial Note: The estimation of national and state-specific adolescent pregnancy and abortion rates was limited by the lack of age-specific abortion and adequate race-specific abortion data for some states. The lack of adequate age-specific abortion data by Hispanic ethnicity in at least half of states for the 3 years also limited this analysis because separate rates for Hispanic and non-Hispanic adolescents could not be computed. State-by-state comparisons of pregnancy rates for whites for states with large Hispanic populations should be interpreted with caution. Moreover, use of abortion data by occurrence rather than by state of residence may have inflated the abortion rate in areas with large metropolitan areas that might draw from adjoining states (e.g., Delaware, DC, and Kansas).

Legally induced abortions reported to CDC may undercount the true number of these abortions (1). Estimates of fetal losses based on NSFG survey data are subject to underreporting because of unrecognized early fetal losses; for females aged <20 years, fetal loss estimates are based on small numbers of pregnancies. Therefore, pregnancy

The District of Columbia is not included in these comparisons because its pregnancy rates were higher than for any state, in part because of large numbers of abortions among nonresidents.

Pregnancy Rates — Continued

		1995	100			1996	96			19	1997		% Change for 15-19-year-olds
Reporting area	<15	15-17	18-19	15-19	<15	15-17	18-19	15-19	<15	15-17	18-19	15-19	from 1995 to 1997
Alabama	10.1	70.8	157.6	105.9	9.7	67.4	156.5	103.6	8.0	63.7	149.3	98.6	6.9-
Alaska					3.0	44.1	116.1	73.4	3.7	39.9	111.1	68.5	
Arizona	6.2	6.69	164.9	107.8	5.7	69.2	156.8	104.3	5.1	62.7	158.1	98.5	7.7-
Arkansas	8.4	68.0	158.3	104.0	8.2	63.9	169.6	105.8	8.8	60.3	165.5	101.8	-2.1
Colorado	4.2	54.1	120.9	80.3	4.2	49.7	122.8	78.2	3,4	48.4	116.9	75.0	9.9-
Connecticut	4.9	53.8	114.6	77.2	5.2	55.7	124.2	81.9	5.3	51.1	122.8	78.6	1.8
Delaware		-						-	11.0	89.0	183.9	127.8	
District of													
Columbia	26.0	* *	*	229.6	22.4	8 8	**	211.2	26.5	176.2	340.8	249.7	8.8
Georgia	11.9	79.2	172.3	115.5	10.5	74.0	169.2	111.4	9.3	71.5	166.1	108.8	-5.9
Hawaii	7.3	58.4	138.7	92.0	4.6	57.8	132.2	89.4	4.7	52.1	124.9	82.6	-10.2
daho	2.9	35.1	105.6	63.4	2.4	34.9	100.7	61.6	2.7	30.8	92.7	99.0	-11.6
Ilinois					7.0		*	100.9	6.1	*	*	92.3	
ndiana	4.8	49.8	131.9	82.3	4.6	48.2	133.5	82.0	3.8	46.4	127.6	78.6	-4.6
Kansas	7.1	59.4	149.9	94.6	5.1	54.6	143.9	89.2	4.9	52.9	143.4	87.9	-7.1
Kentucky	5.8	55.8	137.9	88.5	5,4	51.6	134.6	85.0	4.8	49.3	131.2	82.5	-6.8
onisiana	8.2	63.7	149.7	98.2	9.8	61.4	145.3	95.1	7.7	58.3	143.6	93.0	-5.3
Maine	2.7	36.3	94.2	58.7	1.8	30.8	6.06	54.1	1.2	30.8	92.9	54.9	-6.5
Maryland	7.0	54.9	123.3	81.3	0.9	47.0	112.9	72.5	4.9	41.8	102.5	65.2	-19.8
Massachusetts	4.3	44.2	113.1	71.7	3.9	42.2	109.9	69.2	4.3	41.3	107.1	67.5	-5.8
Michigan	5.2	49.9	127.8	1.08	4.6	46.7	121.3	75.7	4.4	41.6	116.3	71.0	-11.4
Ainnesota	2.7	34.0	92.9	56.4	2.6	33.4	93.6	56.3	2.8	32.0	94.8	6.55	-0.8
Mississippi	10.7	73.4	147.7	103.0	8.6	66.4	144.1	97.5	9.4	64.7	143.3	36.2	-6.7
Missouri	5.2	46.7	130.6	79.1	4.5	44.8	127.0	76.5	4.0	41.5	120.3	72.0	0.6-
Montana	2.7	43.8	118.5	72.8	3.2	40.6	111.5	68.2	3.1	39.4	110.7	67.1	7.7-
Vebraska	3.1	38.8	103.5	64.6	2.9	39.0	109.4	67.0	4.8	37.4	102.3	63.0	-2.4
Jevada	6.7	74.5	185.1	117.1	7.2	69.7	178.0	111.4	5.9	68.9	170.1	107.5	-8.1
New Jersey	5.0	46.2	112.3	71.7	4.7	46.0	109.9	70.7	3.8	39.8	105.8	65.3	6.8
New Mexico	9.9	70.1	164.5	106.8	5.8	69.1	158.3	103.6	5.7	9.99	154.2	100.5	6.5-
Vew York	7.8	6.69	159.8	105.6	8.2	69.2	164.0	107.0	7.0	63.3	151.0	98.1	-7.1
Vorth Carolina	9.6	75.4	168.4	112.3	8.1	68.7	166.0	107.6	7.1	62.1	162.9	102.1	-9.2
<b>Jorth Dakota</b>	11	30.7	8.96	56.3	11	29.8	91.9	53.9	2.4	25.0	84.9	48.2	-14.3
Ohio	4.8	51.5	132.1	83.1	4.3	46.8	127.6	78.5	5.5	48.6	132.5	81.5	-1.9

-10.0	0.0	2 1	15.7	100	000	0.3	-10.8	7.75	0.9-	0.0	23	0 0
64.4	010	89.2	25.5	98.6	108.4	58.0	57.7	77.8	82.8	69.7	57.8	52.8
107.2	148.8	1344	91.2	157.8	169.3	92.6	101.4	122.9	131.4	111.7	94.0	91.6
37.0	53.1	58.5	32.3	59.5	69.3	32.5	30.7	47.0	51.9	40.5	34.9	28.9
4.4	5.1	90	2.7	7.1	6.2	2.6	2.4	80.0	4.9	2.4	3.8	**
67.8	90.5	91.0	56.3	100.6	114.1	58.7	64.5	81.4	85.6	6.69	59.4	54.5
109.4	151.6	135.1	90.2	159.6	175.8	94.0	108.9	126.3	135.2	112.0	8.96	91.1
41.0	50.5	60.7	34.3	62.1	74.2	33.2	36.9	50.6	53.7	41.1	35.7	31.9
5.2	6.2	7.6	2.0	7.9	7.0	2.3	=	5,4	4.5	3.5	3.5	11
71.6	93.4	95.6	59.3	104.4	116.3	57.8	64.7	84.2	88.2	73.2	59.1	58.0
113.8	154.5	141.2	98.5	167.0	176.8	91.9	109.3	127.7	136.9	117.4	95.2	102.7
44.4	54.1	64.6	33.9	64.0	76.6	34.2	36.7	54.4	56.7	43.7	36.1	30.7
5.6	5.3	8.8	1.8	7.7	7,3	2.2	3.4	6.2	5.2	3.7	4.1	11.
Pennsylvania	Rhode Island	South Carolina	South Dakota	Tennessee	Texas	Utah	Vermont	Virginia	Washington	West Virginia	Wisconsin	Wyoming

\* Per 1000 adolescent females in the appropriate age group (13-14 years for < 15-year-old age group).

Abortion data by age for 1995–1997 were not reported for California, Florida, lowa. New Hampshire, and Oklahoma.

Percent changes were computed on the basis of unrounded rates. All but italicized changes in rates were statistically significant at p<0.05.

Pregnancy rates and percent change could not be calculated because the state did not provide abortion data by age for certain years.

\*\* Pregnancy rates could not be calculated because the reporting area did not provide abortion data for certain age groups.

\*\* Pregnancy rate was not calculated for groups with <20 pregnancies or <1000 adolescent females.

TABLE 3. Pregnancy rates\* among adolescents aged 15-19 years and percentage change in rate<sup>1</sup>, by race<sup>1</sup> and state<sup>1</sup> — United States, 1995-1997

	19	95	19	996	19	997		ge in rate 95 to 1997
State	White	Black	White	Black	White	Black	White	Black
Alabama	86.2	145.4	83.8	143.4	81.4	132.0	-5.5	-9.2
Alaska	**	**	61.0	11	泰泰	**	**	**
Arizona	108.7	112.3	105.5	108.8	100.5	109.0	-7.5	-3.0
Arkansas	88.3	155.7	93.0	149.9	89.1	145.0	0.9	-8.0
Colorado	99.	9.9	55	55	73.1	108.1	55	55
Delaware	**	**	**	**	102.7	208.8	**	**
Georgia	90.4	164.7	87.8	156.8	85.4	152.9	-5.5	-7.2
Hawaii	50.7	77.4	52.7	59.4	46.3	65.6	-8.6	-15.2
Idaho	63.3	50	61.1	11	55.7	11	-11.9	11
Indiana	74.0	159.2	73.0	164.1	70.4	155.1	-4.9	-2.5
Kansas	85.8	208.8	81.3	191.4	79.8	194.5	-7.0	-6.9
Kentucky	82.2	154.0	78.9	148.0	77.0	137.8	-6.4	-10.5
Louisiana	71.8	135.59	70.2	130.0%	69.2	125.89	-3.7	-7.199
Maine	57.7	**	53.3	81	54.4	11	-5.7	11
Maryland	58.1	132.4	51.2	120.3	43.8	111.4	-24.6	-15.8
Minnesota	47.1	217.4	47.3	210.6	46.4	213.5	-1.4	-1.8
Mississippi	72.7	137.6	67.8	131.3	68.1	128.6	-6.3	-6.5
Missouri	66.2	161.6	63.6	158.2	60.9	141.4	-8.1	-12.5
Montana	65.4	11	61.5	11	59.2	11	-9.5	11
Nebraska	**	**	**	**	55	55	**	**
Nevada**	117.8	140.9	109.6	145.5	105.8	145.1	-10.2	3.0
New Jersey	46.4	175.6	45.3	169.7	42.0	160.8	-9.5	-8.4
New Mexico	108.6	100.3	106.7	90.5	104.1	86.1	-4.2	-14.2
New York	84.8	190.5	85.0	197.0	78.0	180.7	-8.1	-5.2
North Carolina	92.4	157.6	87.7	152.0	84.4	142.7	-8.7	-9.4
North Dakota	49.4	11	47.2	11	41.5	11	-16.0	11
Ohio	69.2	173.0	65.2	163.1	67.4	173.1	-2.5	0.1
Oregon	90.6	183.2	89.3	191.1	86.0	170.9	-5.0	-6.7
Pennsylvania	53.6	210.0	51.3	195.8	47.8	195.5	-10.8	-6.9
Rhode Island	83.4	206.3	82.7	167.6	82.9	177.6	-0.6	-13.9
South Carolina	78.0	123.4	73.8	118.2	72.9	115.8	-6.6	-6.2
South Dakota	48.3	11	45.8	11	45.0	11	-6.9	11
Tennessee	87.4	169.5	85.3	156.9	82.3	158.0	-5.8	-6.8
Texas	114.3	142.1	111.8	143.0	106.7	132.9	-6.7	-6.5
Utah	56.3	11	57.5	11	56.8	11	0.8	11
Vermont	65.2	11	64.9	11	58.1	**	-10.9	11
Virginia	68.6	138.1	65.5	134.5	62.1	129.3	-9.4	-6.4
Washington	55	150.1	55	15	55	99	99	59
West Virginia	71.1	133.4	68.3	118.1	68.1	116.1	-4.2	-13.0
Wisconsin	46.3	196.8	46.3	197.6	44.9	198.3	-3.0	0.8
Wyoming	**	**	**	**	51.8	11	88	**

\* Per 1000 adolescent females.

Percent changes were computed on the basis of unrounded rates. All but italicized changes in rates were

statistically significant at p<0.05.

1 Pregnancy in Hispanic women is included in rates for white and black adolescents. Race-specific rates, especially for white adolescents, may reflect higher fertility among Hispanic adolescents in states with large Hispanic populations, including Arizona, Colorado, Nevada, New Jersey, New Mexico, New York, Oregon, Rhode Island, Texas, and Washington. Pregnancy rates for adolescents of races other than white or black are not presented because the composition of this category varied widely by state and because abortion information was not available on the race breakdown of "others" for each state.

\*Pregnancy rate and percent change could not be calculated for the following areas because they did not provide

abortion data by age and race for 1995-1997: California, Connecticut, District of Columbia, Florida, Illinois, Iowa,

Massachusetts, Michigan, New Hampshire, and Oklahoma.

\*\* Pregnancy rate and percent change could not be calculated because state did not provide abortion data by age and race for certain years.

"Pregnancy rate and percent change could not be calculated for age groups with <20 pregnancies or <1000 adolescent females.

\*\* Pregnancy rate and percent change could not be calculated because age or race information was missing for >15% of women who had an abortion.

11 Rate and percent change is for all races other than white.

totals based on births, legally induced abortions reported to CDC, and fetal loss estimates may underestimate the actual pregnancy rate. However, underreporting likely remains relatively constant from year to year and is therefore unlikely to affect the trends shown in this report.

Sexual experience, sexual activity, and effective contraceptive use are important determinants of changes in pregnancy rates. The decline in pregnancy rates among females aged 15–19 years have been attributed to stable rates of sexual experience and activity among this group and increased use of condoms (4,8). Increased use of long-acting hormonal methods introduced in the early 1990s also has been associated with the decline (9).

Sustaining the downward trend in adolescent pregnancy will require addressing complex individual and community-level factors that can affect adolescents' sexual and reproductive behavior. Community- and school-based programs designed to reduce adolescent pregnancy that address risk factors and specific skills to postpone sexual experience and increase contraceptive use may be more effective in reducing adolescent pregnancy than programs focusing exclusively on changing sexual beliefs or behavior (10). Effective programs also include strong educational components, messages targeting different groups of adolescents, and youth development approaches that will strengthen self-esteem and planning for the future (10). Scientific evaluation of adolescent pregnancy prevention measures is an essential component of these community-based programs. The identification of effective strategies will assist state and local agencies in implementing successful approaches to continuing the downward trend in adolescent pregnancy.

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# Silicosis Screening in Surface Coal Miners — Pennsylvania, 1996-1997

Silicosis is an occupational respiratory disease caused by inhaling respirable crystal-line silica dust. Silicosis is irreversible, often progressive (even after exposure has ceased), and potentially fatal. Exposure to silica dust occurs in many occupations, including mining (1). During 1996–1997, surface coal miners at eight sites in Pennsylvania were screened to estimate the prevalence of silicosis, to identify risk factors for silicosis, and to refer miners with a possible diagnosis of silicosis or other conditions for medical evaluation and treatment. This report summarizes the results of the screening, which indicated that an increased prevalence of and risk for silicosis is associated with miners' age and years of drilling experience, and provides recommendations for preventing silicosis among miners.

Enrollment in the screening was voluntary and available to anyone who had worked in surface mining for 1 year or more. Surface miners were informed of the screening by Mine Safety and Health Administration (MSHA) inspectors during routine mine visits. Screening was performed by a multiagency team from the Chronic Respiratory Disease Program of the Pennsylvania Department of Health (team leader), the Department of Health Evaluation Sciences of the Pennsylvania State University College of Medicine, MSHA, and CDC's National Institute for Occupational Safety and Health (NIOSH).

Screening was conducted during May–June 1996, at five mine sites in bituminous coalfields in western Pennsylvania (Altoona, Clearfield, Farmington, Indiana, and Somerset) and, in June 1997, at three mine sites in anthracite coalfields in eastern Pennsylvania (Centralia, Pottsville, and Wilkes-Barre). The screening was divided by coal type and region because of differences in geology and mining practices. Screening consisted of anterior-posterior chest radiographs; spirometry; and a survey containing questions about demographics; medical, work, and smoking history; and workplace exposures. Silicosis was defined as a radiographic finding of International Labour Organization (ILO) classification of profusion category ≥1/0 (2); classification was based on consensus of at least two of three NIOSH-certified B readers\*. Descriptive analyses were performed on all variables collected from the radiographs, spirometry, and surveys. Multivariable logistic regression models were used to determine risk factors for developing silicosis.

During 1996–1997, 1250 current and former coal miners were screened at the eight sites (664 in western and 586 in eastern Pennsylvania); and data from 1236 miners were suitable for analysis (Figure 1). Screened miners were almost exclusively white (99.9%), male (99.5%), and non-Hispanic (97.6%); the mean age was 46.2 years (range: 18–87 years). Of 1221 miners, 289 (23.7%) were current smokers, and 729 (59.7%) had ever smoked: 1120 (90.7%) of 1235¹ were employed full-time.

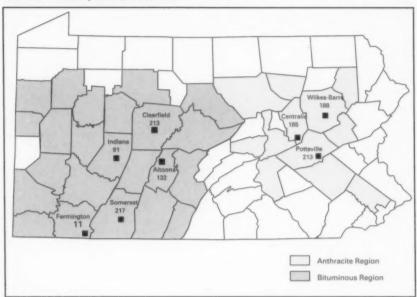
Radiographic evidence of silicosis was found in 83 (6.7%) of 1236 screened miners. Prevalence of silicosis did not vary by smoking status, and no significant differences in prevalence were noted by site except among the 213 participants at Clearfield (16.0%; p=0.001). When data from Clearfield were excluded, the prevalence of silicosis was

<sup>\*</sup>A physician certified by NIOSH as having competency in the classification of chest radiographs to detect pneumoconiosis using ILO guidelines. If at least two of the three B readers categorized the profusion as ≥1/0, the miner was classified as having silicosis; if at least two readers indicated the film was negative (<1/0), the miner was classified as not having silicosis.

<sup>&#</sup>x27;Denominators vary because of nonresponse to specific questions.

Silicosis - Continued

FIGURE 1. Number of surface coal miners participating in silicosis screening, by region and site — Pennsylvania, 1996–1997



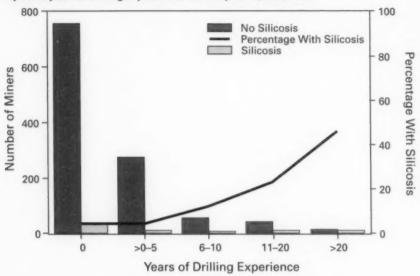
similar for western (5.2%) and eastern (4.5%) Pennsylvania. The odds ratio (OR) for silicosis at the Clearfield site compared with other western sites was 4.4 (95% confidence interval [CI]=2.3–8.5); the OR for silicosis at eastern sites was 1.1 (95% CI=0.6–2.1) compared with western sites, excluding Clearfield.

The mean age of miners with silicosis was 50.6 years and without silicosis was 45.6 years (p=0.0002). When age was modeled as a quadratic function in the logistic regression model, a significant increase in silicosis occurred with increasing age. Compared with miners aged 30 years, miners aged 40, 50, and 60 years had progressively increased odds of silicosis: 3.7 (95% Cl=1.7–8.2), 7.8 (95% Cl=2.4–25.3), and 9.7 (95% Cl=3.0–31.6), respectively. Silicosis prevalence increased as the number of reported years of drilling experience increased, from 37 (4.7%) of 792 miners reporting no drilling experience to 12 (46%) of 26 in miners reporting >20 years experience (Figure 2). Compared with miners with ≤5 years of drilling experience, those who had 6–10, 11–20, and >20 years experience had progressively increased odds of silicosis: 4.3 (95% Cl=1.6–11.8), 7.0 (95% Cl=2.6–18.6), and 14.5 (95% Cl=4.8–43.6), respectively.

Reported by: PA Tyson, MSW, Chronic Respiratory Disease Program, Pennsylvania Dept of Health. JL Stauffer, MD, EA Mauger, PhD, JE Caulfield, MS, Pennsylvania State Univ College of Medicine, Hershey. DW Conrad, KG Stricklin, Mine Safety and Health Administration, US Dept of Labor. Div of Respiratory Disease Studies; Office for Mine Safety and Health, Pittsburgh Research Laboratory, National Institute for Occupational Safety and Health, CDC.

Silicosis - Continued

FIGURE 2. Number and percentage of surface coal miners with and without silicosis, by reported years of drilling experience — Pennsylvania, 1996–1997



**Editorial Note:** This report underscores the risk for silicosis associated with surface coal mining operations. Previous studies identified an increased risk for silicosis among rock drillers (3), and this report corroborates the increased prevalence of silicosis among surface coal mining drillers (4.5).

The findings in this report are subject to at least four limitations. First, the sample was voluntary and represented approximately 40% of Pennsylvania surface coal miners (63% of anthracite miners and 29% of bituminous miners<sup>5</sup>); therefore, the results do not necessarily represent all surface coal miners in Pennsylvania or the United States. Silicosis prevalence may be underestimated if miners with confirmed or suspected silicosis did not participate or may be overestimated if a higher percentage of affected workers participated. Second, B reader variability in the interpretation of chest radiographs was a possibility, although the study methods were designed to limit the effects of reader variability (6). Third, prevalence differences across sites must be interpreted cautiously; miners were not restricted in their choice of screening site, and the latency period for silicosis, several years to several decades following exposure, makes it difficult to determine a specific source of exposure in workers. Finally, the study collected data on years of drilling experience and not on years of overall mining experience; some miners may have performed numerous duties at different mines throughout the region.

Data from MSHA, part 50. The Federal Mine Safety and Health Act of 1977 requires all mine operators to record and report to MSHA data on occupational injuries; illnesses; certain noninjurious accidents; and related employment, work time, and operating activity information. MSHA data cited for number of employees are for mine operator employees only; state-specific data are not available for contractor employees.

Silicosis - Continued

According to MSHA part 50 data for 1996 and 1997, Pennsylvania had almost one third of the U.S. surface coal mines; however, because of the many small operations in Pennsylvania, 3205 (approximately 10%) of the 31,308 surface coal miners in the United States are employed in Pennsylvania. Smaller mine operations may lack resources required to purchase or maintain optimal dust-control equipment, and small-scale operations represent special challenges for enforcement activities. The reason for the higher silicosis prevalence at Clearfield is unknown; however, influences may include site-specific geologic factors (e.g., quartz content of overlying rock), past work practices, mining methods, types of controls, or machinery maintenance.

Because no effective treatment exists for silicosis, prevention through exposure control is essential (7). When proper practices are not followed or controls are not maintained, silica exposures can exceed the MSHA Permissible Exposure Limit (PEL) or the NIOSH Recommended Exposure Limit (REL)\*. Effective engineering controls in the mining environment include dust suppression (e.g., wet drilling), dust collection (e.g., dry drilling with particulates exhausted through a dust collection system), and use of enclosed isolation systems (e.g., air conditioned cabs under positive pressure and equipped with both filtered air supply and filtered recirculated air). In 1994, MSHA and NIOSH implemented a regional, and later a national, silicosis prevention program that promoted educational efforts and allowed coal mine inspectors to issue citations on the basis of visual inspection of dust-producing drilling equipment rather than more extensive sampling (8). As a result, some high-risk equipment has been discarded by the mines; however, because of the long latency period that usually precedes clinical onset of silicosis, the impact of these reforms on the incidence of silicosis remains unclear.

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<sup>&</sup>lt;sup>1</sup>PEL for coal mining is 2.0 mg of respirable coal dust per cubic meter of air (2 mg/m³); however, when the respirable quartz (crystalline silica) content of the dust exceeds 5%, a reduced PEL is computed by dividing the percentage of respirable quartz into the number 10 (Title 30, Code of Federal Regulations, Part 71.101). NIOSH-recommended exposure limit (REL) for respirable crystalline silica is a 10-hour, time-weighted average level of 50 µg/m³.

# Hepatitis B Vaccination Coverage Among Asian and Pacific Islander Children — United States, 1998

Asian and Pacific Islander (API) children in the United States have high rates of hepatitis B virus (HBV) infection (1-3). To prevent these infections, hepatitis B vaccination has been recommended for these children since the vaccine was first licensed by the Food and Drug Administration in 1981 (4). Recommendations have included universal hepatitis B vaccination of API infants beginning in 1990 and catch-up vaccination for API children aged <7 years (5). These recommendations were reinforced in 1991 when hepatitis B vaccination was recommended for all infants, particularly in populations such as API children with high rates of early childhood HBV infection (6). In 1995, vaccination was recommended for unvaccinated API children aged <11 years and catch-up vaccination for children aged 11-12 years who had not received hepatitis B vaccine (HepB) (7). Series completion among API children aged 19-35 months increased from 39% in 1994 to 88% in 1997 (8). However, among older API children, hepatitis B vaccination coverage was 10% in 1995 (7). In 1998, to examine trends in hepatitis B vaccination catch-up coverage among API children born before 1994, surveys were conducted in six U.S. cities. This report summarizes the results of the surveys, which indicate varying coverage among API children and suggest a need for continued focused vaccination programs for this population.

In three cities (Milwaukee, St. Paul, and Seattle), state or county health departments evaluated hepatitis B vaccination programs conducted in the API community. These cities were compared with three other cities (Dallas, Houston, and Washington, D.C.) where no vaccination programs were targeted for API children. Sampling and data collection methods varied between cities. In Milwaukee, students aged 5-14 years with Lao/ Hmong surnames were sampled randomly from all public schools. In St. Paul and Seattle, children aged 4-13 and 4-14 years, respectively, were selected using a populationbased cluster sample technique. In these three cities, parents or guardians were interviewed in person, and vaccination data were validated with written records from parents, schools, or health-care providers. In Dallas, Houston, and Washington, D.C., a random sample of persons with Vietnamese surnames was selected from area telephone directories, primary caretakers of persons aged 3-18 years were interviewed by telephone, and health-care providers were contacted to verify vaccination histories. For each city, a range of the percentage vaccinated was estimated using the total sample to represent the lower percentage and only those children identified with health-care providers for the higher percentage (Table 1).

Children born during 1984–1993 were included for this report. The third HepB dose (HepB-3) was counted if it had been administered at or after age 164 days with an interval of at least 108 days between doses 1 and 3. LOWESS plots (9) were constructed to compare city-specific trends in yearly HepB administration rates among children eligible for vaccination. The percentage of all children in each survey who completed the HepB series by 1998 ranged from 14% (Houston) to 67% (St. Paul) (Table 1). For all children in all surveys combined, the weighted average HepB series completion rate was 42% and did not differ by place of birth (born in the United States versus foreign-born; n=1232 and 708, respectively). Series completion rates at or before age 12, 18, and 60 months were 12%, 16%, and 30%, respectively.

Hepatitis B Vaccination Coverage - Continued

TABLE 1. Hepatitis B vaccination coverage rates among Asian and Pacific Islander children born during 1984-1993 - six-city survey, 1998

		Hous	eholds		No.	No.		% of	% of
City	Total	Response rate	No. eligibles identified	Eligible response rate	children in sample	children with providers	Ethnicity	children receiving Hep B-1*	children receiving Hep B-31
Without program									
Dallas <sup>6</sup>	3801	72%	549	91%	332	(177)9	Vietnamese	28% (52%)9	18% (36%)
Houston <sup>5</sup>	4743	65%	539	94%	314	(132)9	Vietnamese	25% (61%)9	14% (36%)
Washington, DC1	3550	79%	503	93%	346	(127)9	Vietnamese	25% (56%)4	15% (43%)
With program									
Milwaukee**	275	99%	271	76%	207		Lao/Hmong	82%	51%
St. Paul <sup>11</sup>	1391	56%	209	96%	586		Hmong	80%	67%
Seattle <sup>11</sup>	4200	95%	272	100%	412		Pan-Asian <sup>18</sup>	79%	65%

\* First dose in the hepatitis B vaccination series.

1 Third dose in the hepatitis B vaccination series. Systematic random sample from telephone directory list of Vietnamese surnames.

1 To be included in this subsample, health-care providers had to report child as patient.

\*\* Systematic random sample from school enrollment list of Lao/Hmong surnames.

"Cluster sample of households within geographic area.
"Primarily Vietnamese (32%), Chinese (19%), Filipino (19%), and Cambodian (12%).

Vaccination coverage was examined separately for cities with and without hepatitis B vaccination programs for API children. HepB coverage was 41%-61% and 2%-11% for cities with and without these programs for the 1980s birth cohorts and increased with the 1990s birth cohorts (Figure 1). The increase was greatest in cities with a designated API vaccination program; combined vaccination coverage in the 1993 birth cohort was 83%. The effect of the 1990 and 1991 recommendations for infant vaccination was observed when vaccination coverage was stratified by age at HepB series completion (Figure 1). In each birth cohort, the proportion of children who completed the HepB series by age 18 months (infant vaccination) increased substantially in cities with and without ongoing programs; however, during 1992-1998, annual catch-up vaccination rates as measured by HepB-3 completion remained 7%-11% per year in cities with ongoing API hepatitis B vaccination programs compared with 0.7%-2.6% per year in cities without such pro-

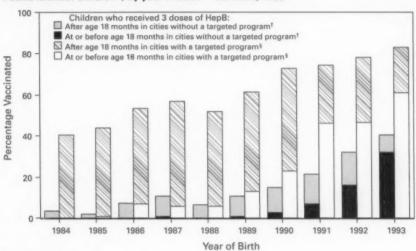
Reported by: C Jenkins, Univ of California, San Francisco, California, M Roddy, Minnesota Dept of Health. L Stewart, Seattle-King County Public Health, Seattle, Washington. M Hurie, Wisconsin Div of Health. J Millen, Association of Schools of Public Health, Atlanta, Georgia. Adult Vaccine-Preventable Diseases Br, Epidemiology and Surveillance Div, National Immunization Program, CDC.

Editorial Note: The findings in this report document the impact of targeted vaccination programs for populations at high risk for childhood HBV infection. In the three cities with ongoing API hepatitis B vaccination programs, coverage increased with each successive birth cohort over a 10-year period, reaching 83% among children born in 1993; however, in cities without programs, the overall vaccination coverage remained low, although coverage also increased with each successive birth cohort.

The three cities with API hepatitis B vaccination programs used three approaches to achieve their higher rates of coverage. In Milwaukee, an active refugee health hepatitis B vaccination program, which began in 1984, provided screening and vaccination services to all new API arrivals. In St. Paul, during the 1989-1991 measles epidemic that resulted in three deaths among Hmong children, a coalition was formed that conducted

Hepatitis B Vaccination Coverage - Continued

FIGURE 1. Coverage with three doses of hepatitis B vaccine (HepB) among Asian and Pacific Islander children\*, by year of birth — six cities, 1998



<sup>\*</sup>n=2197.

a multimedia health-promotion campaign and health-care provider education and outreach. In Seattle, state and local health departments and two clinics that served a large proportion of the API population educated providers about the need for catch-up coverage among API children and, during 1996–1997, conducted a 16-month, citywide middle school and high school-based hepatitis B vaccination program for all public school students.

The findings in this report are subject to at least three limitations. First, the populations studied may not represent the nationwide API population. Second, the cities with and without programs may not represent all U.S. cities with and without targeted catchup hepatitis B vaccination programs. Third, biases may have resulted from a loss of randomness in sampling, nonrespondents whose vaccination rates differ from respondents, and missing vaccination records that may have caused inaccurate coverage estimates.

Data from the six city surveys and from other cities and state reports (10) indicate that 40% of all API children in the United States aged 7–18 years have completed their HepB series (CDC, unpublished data, 1999). This low rate of coverage in a group at risk for HBV infection underscores the need for increased efforts to continue to provide catch-up vaccination to these children. In addition, catch-up efforts among preschool children are needed in communities where many API infants do not receive HepB. API children born since 1988 who were not vaccinated as infants or caught up as young

Dallas, Houston, and Washington, D.C.

<sup>&</sup>lt;sup>1</sup> Milwaukee, St. Paul, and Seattle.

Hepatitis B Vaccination Coverage - Continued

children should be vaccinated routinely at age 11–12 years. Although eventual high rates of universal adolescent vaccination can be expected for API children living in the 20 states and Washington, D.C., with existing middle school entry laws, special efforts will be needed to ensure vaccination of API children in states without such laws. Because no established vaccination visits exist for older adolescents, hepatitis B vaccination will depend primarily on self-identification, community-based programs, and health-care providers who are aware of the high risk for HBV infection among API children and who can meet specific API cultural and language needs (CDC, unpublished data, 1999). Community-based catch-up hepatitis B vaccination programs have been the mission of the National Task Force on Hepatitis B Immunization, Focus on Asians and Pacific Islanders (on the World-Wide Web at http://aapihp.com\*). Successful catch-up initiatives to protect API children should be implemented as quickly as possible.

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# Notice to Readers

# Delayed Supply of Influenza Vaccine and Adjunct ACIP Influenza Vaccine Recommendations for the 2000–01 Influenza Season

Annual vaccination against influenza is the primary means for minimizing serious adverse outcomes from influenza virus infections. These infections result in approximately 20,000 deaths and 110,000 hospitalizations per year in the United States (1). The

<sup>\*</sup>References to sites of non-CDC organizations on the World-Wide Web are provided as a service to *MMWR* readers and do not constitute or imply endorsement of these organizations or their programs by CDC or the U.S. Department of Health and Human Services. CDC is not responsible for the content of pages found at these sites.

amount of trivalent inactivated influenza vaccine produced for distribution in the United States has increased substantially; in 1999, four manufacturers produced a combined total of 80 to 85 million doses.

For the 2000–01 influenza season in the United States, lower than anticipated production yields for this year's influenza A(H3N2) vaccine component and other manufacturing problems are expected to lead to a substantial delay in the distribution of influenza vaccine and possibly substantially fewer total doses of vaccine for distribution than last year. A more precise estimate of the vaccine supply will be available as production progresses during the summer. Because many vaccine providers currently are planning their fall vaccination activities, CDC and the Advisory Committee on Immunization Practices (ACIP) are issuing adjunct influenza vaccination recommendations beyond those made by ACIP on April 14, 2000 (1). The adjunct recommendations are specific to the 2000–01 influenza season.

### Adjunct Influenza Vaccine Use Recommendations for the 2000-01 Influenza Season

- Implementation of organized influenza vaccination campaigns should be delayed.
  Health-care providers, health organizations, commercial companies, and other
  organizations planning organized influenza vaccination campaigns for the 2000–01
  influenza season should delay vaccination campaigns until early to mid-November.
  The purpose of this recommendation is to minimize cancellations of vaccine campaigns
  and wastage of vaccine doses resulting from delays in vaccine delivery.
- 2. Influenza vaccination of persons at high risk for complications from influenza and their close contacts should proceed routinely during regular health-care visits. Routine influenza vaccination activities in clinics, offices, hospitals, nursing homes, and other health-care settings (especially vaccination of persons at high risk for complications from influenza, health-care staff, and other persons in close contact with persons at high risk for complications from influenza) should proceed as normal with available vaccine.
- 3. Provider-specific contingency plans for an influenza vaccine shortage should be developed. All influenza vaccine providers, including health-care systems and organizers of vaccination campaigns, should develop a provider-specific contingency plan to maximize vaccination of high-risk persons and health-care workers. These plans should be available for implementation if a vaccine shortage develops.

#### Use of Influenza Antiviral Medications

There are no new recommendations for the use of influenza antiviral drugs. The approved usage (i.e., for treatment or chemoprophylaxis), age group, dosage, route of administration, metabolism, and adverse reactions of these agents vary (1), and all of them require prescription by a physician. Influenza antiviral drugs are useful for controlling influenza outbreaks in specific and circumscribed situations, such as nursing homes. In addition, long-term antiviral chemoprophylaxis of high-risk institutionalized residents or some persons at high risk for complications from influenza might be indicated if vaccine either is unavailable, ineffective (e.g., severely immunocompromised persons), or contraindicated.

However, these drugs are not a substitute for influenza vaccine. Even if an influenza vaccine shortage develops, CDC and ACIP do not support their routine and widespread use as chemoprophylaxis against influenza because this is an untested and expensive strategy that could result in large numbers of persons experiencing adverse effects.

### **Additional Discussion**

In the United States, 70 to 76 million persons (approximately 35 million persons aged ≥65 years; 33 to 39 million persons aged <65 years with high-risk medical conditions; and 2 million pregnant women) are at high risk for serious complications from influenza infections, including hospitalizations and deaths. The expected delay in influenza vaccine distribution and a possible shortage for the 2000–01 influenza season has raised difficult questions of how to maximize protection against influenza for these persons. One complicating factor is that many vaccine providers must plan their fall vaccination activities now even though the vaccine supply is uncertain. Given the current situation, CDC and ACIP have issued modified recommendations for the 2000–01 season emphasizing the delay of organized influenza vaccine campaigns until November, the continuation of routine vaccination activities during regular health-care visits, and the development of provider-specific contingency plans in case a vaccine shortage should develop. There are additional important points worth emphasizing in addition to these main recommendations:

- Influenza vaccine administered after mid-November can still provide substantial protective benefits. In general, ACIP recommends that routine vaccination of persons at high risk for complications from influenza begin in September. In previous years, ACIP has recommended that organized campaigns take place during October through mid-November. These timing recommendations balance several considerations, including the desirability of administering vaccine before substantial seasonal influenza activity has begun but not vaccinating so early such that vaccine antibody titers might substantially decrease in some persons. Nonetheless, many persons who should receive influenza vaccine remain unvaccinated after mid-November, and for many of these persons, influenza vaccination after mid-November will be beneficial. For the 2000–01 season, it is particularly important for vaccine providers to continue to administer vaccine after mid-November.
- Once vaccine is available, health-care workers should provide vaccine to persons at high risk for complications from influenza as is normally done. This is particularly important for young children at high risk who are receiving influenza vaccination for the first time and who require two doses of vaccine.
- Minimizing wastage of influenza vaccine is important. In particular, influenza vaccine purchasers should refrain from placing duplicate orders with multiple companies to minimize the amount of vaccine that is returned to a manufacturer and discarded. Options to promote redistribution of vaccine that otherwise would be returned or discarded are being developed.
- In 2000, ACIP broadened its influenza vaccine recommendations to include all
  persons aged 50–64 years. This recommendation was based, in part, on an effort
  to increase vaccination coverage of persons in this age group with high-risk
  conditions. In the context of a possible vaccine shortage, it would be appropriate
  for contingency plans covering this age group to focus primarily on vaccinating
  persons with high-risk conditions rather than this entire age group.
- Influenza vaccine is routinely recommended for persons in close contact with
  persons at high risk for complications from influenza because such persons are in
  a position to transmit influenza virus infection to high-risk persons. Vaccination of
  health-care workers has been highlighted in particular because health-care workers have frequent and close contact with many different high-risk persons at a
  time when high-risk persons are particularly vulnerable.

As new information becomes available, CDC and the Food and Drug Administration (FDA) will issue updates. In the meantime, ACIP and CDC request that persons and organizations planning to administer influenza vaccine, as well as members of the general public, join in these efforts to maximize protection of persons most likely to develop serious and life-threatening complications from influenza. FDA, CDC, ACIP, National Institutes of Health, and vaccine manufacturers will continue to work together to facilitate the availability of influenza vaccine for the upcoming season and to minimize the adverse impact of an influenza vaccine shortage if one should develop. If a substantial vaccine shortage appears imminent, or if the situation warrants, then CDC and ACIP will issue further recommendations.

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 CDC. Prevention and control of influenza: recommendations of the Advisory Committee on Immunization Practices (ACIP). MMWR 2000;49(no. RR-3).

## Notice to Readers

# Summary of the Joint Statement on Thimerosal in Vaccines

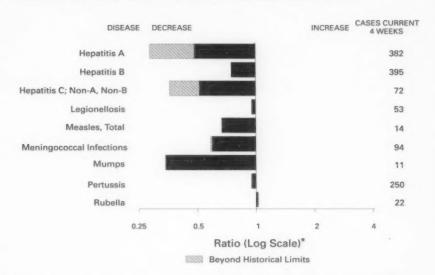
In June 2000, a joint statement on thimerosal\* in vaccines was prepared by the American Academy of Family Physicians (AAFP), the American Academy of Pediatrics (AAP), the Advisory Committee on Immunization Practices (ACIP), and the Public Health Service (PHS) in response to 1) the progress in achieving the national goal declared in July 1999 to remove thimerosal from vaccines in the recommended childhood vaccination schedule, and 2) results of recent studies that examined potential associations between exposure to mercury in thimerosal-containing vaccines and health effects. In this statement, AAFP, AAP, ACIP, and PHS recommend continuation of the current policy of moving rapidly to vaccines that are free of thimerosal as a preservative. Until adequate supplies are available, use of vaccines that contain thimerosal as a preservative is acceptable.

A joint statement issued by AAP and PHS in July 1999 and agreed to by the AAFP later in 1999 established the goal of removing thimerosal as soon as possible from vaccines routinely recommended for infants. The goal was established as a precautionary measure. No evidence existed of any harm caused by low levels of thimerosal in vaccines. Public concern had been expressed about the health effects of mercury exposure of any sort, and the elimination of mercury from vaccines was considered a feasible means of reducing an infant's total exposure to mercury in a world where other environmental sources of exposure are more difficult or impossible to eliminate (e.g., certain foods).

(Continued on page 631)

<sup>\*</sup>Thimerosal is a derivative of ethylmercury and has been used as an additive to biologics and vaccines since the 1930s because it is effective in killing bacteria and in preventing bacterial contamination, particularly in opened, multidose containers. The full text of this statement is available on the World-Wide Web at http://www.aap.org/policy/camp/20.html, http://www.aap.org/policy/jointthim.html, and http://wwww.cdc.gov/nip/vacsafe/concerns/thimerosal/joint\_statement\_00.htm. References to sites of non-CDC organizations on the World-Wide Web are provided as a service to MMWR readers and do not constitute or imply endorsement of these organizations or their programs by CDC or the U.S. Department of Health and Human Services. CDC is not responsible for the content of pages found at these sites.

FIGURE I. Selected notifiable disease reports, United States, comparison of provisional 4-week totals ending July 8, 2000, with historical data



\*Ratio of current 4-week total to mean of 15 4-week totals (from previous, comparable, and subsequent 4-week periods for the past 5 years). The point where the hatched area begins is based on the mean and two standard deviations of these 4-week totals.

TABLE I. Summary of provisional cases of selected notifiable diseases, United States, cumulative, week ending July 8, 2000 (27th Week)

		Cum. 2000		Cum. 2000
Anthrax			HIV infection, pediatric*1	108
Brucellosis*		26	Plague	4
Cholera			Poliomyelitis, paralytic	1
Congenital ru	bella syndrome	4	Psittacosis*	8
Cyclosporiasis		16	Rabies, human	
Diphtheria			Rocky Mountain spotted fever (RMSF)	128
Encephalitis:	California serogroup viral*	4	Streptococcal disease, invasive, group A	1,671
	eastern equine*		Streptococcal toxic-shock syndrome*	55
	St. Louis*		Syphilis, congenital <sup>5</sup>	67
	western equine*		Tetanus	12
Ehrlichiosis	human granulocytic (HGE)*	48	Toxic-shock syndrome	86
	human monocytic (HME)*	20	Trichinosis	4
Hansen disea	se (leprosy)*	20 30	Typhoid fever	162
Hantavirus pu	Ilmonary syndrome*1	13 46	Yellow fever	
Hemolytic ure	emic syndrome, postdiarrheal*	46		

<sup>&</sup>quot;Not notifiable in all states.

<sup>&</sup>quot;Updated weekly from reports to the Division of Viral and Rickettsial Diseases, National Center for Infectious Diseases (NCID).

\*Updated monthly from reports to the Division of HIV/AIDS Prevention — Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention (NCHSTP). Last update June 25, 2000.

\*Updated from reports to the Division of STD Prevention, NCHSTP.

TABLE II. Provisional cases of selected notifiable diseases, United States, weeks ending July 8, 2000, and July 10, 1999 (27th Week)

	All	ne	Chlan	to the state of	Comton	and diamin	NET		coli O157:H	
Demostre Acce	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.	Cum.
Reporting Area UNITED STATES	2000° 20,482	1999 22,981	2000 294,973	1999 334,806	2000 632	<b>1999</b> 912	1,151	1999 852	<b>2000</b> 676	1999 873
NEW ENGLAND Maine N.H.	1,213 16 18 11	1,109 29 30 6	11,011 694 511 276	10,893 566 512 247	34 9 4 13	46 9 5 6	122 7 10	127 11 14 15	104 6 9 4	120 16 8
Mass. R.I. Conn.	776 49 343	702 63 279	5,128 1,211 3,191	4,605 1,216 3,747	6 2	23	54 8 38	58 6 23	50 5 30	56 7 33
MID. ATLANTIC Jpstate N.Y. N.Y. City N.J. Pa.	4,928 572 2,620 1,036 700	5,893 727 2,995 1,146 1,025	20,385 N 5,527 3,553 11,305	35,269 N 14,943 6,319 14,007	64 39 7 7	193 57 113 15	128 103 7 18 N	59 38 4 17 N	67 43 16 8	66 5 60 1
E.N. CENTRAL Ohio nd. III. Mich. Wis.	2,052 306 191 1,198 255 102	1,498 246 189 677 307 79	48,339 12,299 6,086 13,112 12,130 4,712	55,283 14,147 6,163 16,539 11,264 7,170	137 23 12 7 28 67	158 20 9 31 22 76	198 42 36 55 40 26	162 54 19 61 28 N	93 25 23 26 19	150 48 20 39 19 24
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr. Kans.	480 87 52 223 1 4 31 82	502 82 52 231 4 11 37 85	17,057 3,282 2,101 6,331 282 865 1,548 2,648	19,494 3,912 2,274 7,097 459 821 1,720 3,211	57 11 17 10 5 5 7	54 13 12 10 4 3 11	182 52 38 48 8 10 15	145 36 27 14 3 5 48	125 49 10 41 8 3 9	184 60 15 24 5 17 61 2
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga.	5,443 94 602 388 385 33 334 434 607 2,566	6,282 80 720 239 335 31 394 579 967 2,947	62,427 1,480 6,384 1,731 7,782 753 11,454 4,880 11,830 16,133	71,896 1,417 6,571 N 7,707 916 11,724 9,166 18,357 16,038	123 4 7 7 7 4 3 11 61 26	165 7 7 7 10 4 86 51	98 12 20 7 19 6 13 21	99 4 7 28 4 22 12 7 15	51 U 15 3 11 2 10 9	86 U 29 27 9 U 19
E.S. CENTRAL Ky. Tenn. Ala. Miss.	1,005 114 407 262 222	1,028 151 402 255 220	23,692 4,167 7,319 7,432 4,774	22,598 3,985 7,109 5,346 6,158	25 1 7 10 7	12 4 4 2 2	50 18 20 5 7	59 14 25 14 6	26 12 12 2	45 11 18 14 2
W.S. CENTRAL Ark. La. Okla. Tex.	1,868 103 336 156 1,273	2,475 90 464 71 1,850	44,100 2,717 9,808 4,019 27,556	45,747 3,082 7,279 4,105 31,281	28 1 8 4 15	21 2 17	68 33 4 9 22	43 5 5 7 26	59 3 18 7 31	53 5 7 6 35
MOUNTAIN MORT. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	755 9 13 6 157 86 244 67 173	852 4 12 3 171 46 422 80 114	18,904 803 966 377 5,582 2,345 5,851 1,240 1,740	18,079 654 869 382 4,186 2,731 6,570 1,092 1,595	88 3 3 12 2 3 9 2	41 7 3 4 17 7 N 3	130 15 14 7 56 5 25 7	67 4 2 3 25 4 11 15 3	70 2 30 3 18 17	60 6 5 15 2 7 19 6
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	2,738 285 89 2,275 10 79	3,342 185 87 3,011 13 46	49,058 6,645 2,626 37,476 1,182 1,129	55,547 6,124 3,251 43,589 946 1,637	122 N 8 114	203 N 76 127	175 55 29 82 2	91 31 22 33	81 43 31	109 40 22 42
Guam P.R. V.I. Amer. Samoa C.N.M.I.	13 518 21	737 15	298	233 U U U	:	UUU	N 4	N 5 U U U U	0000	0000

N: Not notifiable. U: Unavailable. ... No reported cases. C.N.M.I.: Commonwealth of Northern Mariana Islands.

\*Individual cases can be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Heath Laboratory Information System (PHLIS).

\*Chlamydia refers to genital infections caused by C. trachomatis. Totals reported to the Division of STD Prevention, NCHSTP.

\*Updated monthly from reports to the Division of HIV/AIDS Prevention — Surveillance and Epidemiology, National Center for HIV, STD, and TB Prevention. Last update June 25, 2000.

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending July 8, 2000, and July 10, 1999 (27th Week)

	Gono	rrhea		rtitis C; Non-B	Legion	nellosis		/me sease
Reporting Area	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum 1999
UNITED STATES NEW ENGLAND Maine N.H. vt. Mass. R.I.	3,007 41 53 30 1,342 304	178,480 3,280 27 49 28 1,281 313	1,275 27 1 3 20 3	1,959 9 1	366 23 2 2 2 2 9 3	454 29 3 3 4 10 3	2,952 734 36 4 275	4,570 1,382 1 1 1 2 372 99
Conn. MID. ATLANTIC Upstate N.Y. N.Y. City N.J. Pa.	1,237 12,964 3,380 2,322 1,958 5,304	1,582 20,325 3,077 7,102 3,759 6,387	36 36	69 34 	73 32 4 37	6 111 27 14 11 59	363 1,690 821 4 287 578	907 2,271 965 67 564 675
E.N. CENTRAL Ohio Ind. III. Mich. Wis.	29,959 7,254 2,808 9,123 9,045 1,729	34,401 8,863 3,294 11,199 7,874 3,171	115 4 1 7 103	1,086 1 1 31 463 590	89 38 20 8 17 6	143 44 18 19 35	59 25 10 1	317 22 14 11 8 262
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr. Kans.	7,460 1,334 465 3,811 6 128 562 1,154	8,228 1,439 529 4,029 43 80 786	345 5 1 314	94 2 90	27 1 4 17	24 1 7 11 1	75 24 4 14	66 13 8 30 1
S. ATLANTIC Del. Md. D.C. Va. W. Va. N. C. S.C. Ga. Fla.	45,123 830 4,221 1,235 4,955 227 9,315 5,783 7,369 11,188	52,805 863 5,881 1,862 5,189 309 10,062 5,227 12,065 11,347	62 6 2 1 9 13 1 2 28	107 29 10 13 24 12 1	80 4 25 1 11 N 8 2 4	55 6 10 13 N 8 7	33 325 34 207 1 49 8 11 2	7 407 36 288 1 29 10 34 3
E.S. CENTRAL Ky. Tenn. Ala. Miss.	17,029 1,740 5,695 5,843 3,751	17,693 1,740 5,644 4,834 5,475	229 17 57 7 148	159 9 54 1 95	11 5 4 2	22 10 10 2	14 3 8 2	40 5 15 10
W.S. CENTRAL Ark. La. Okla. Tex.	22,989 1,427 6,894 1,731 12,937	25,753 1,529 5,987 2,083 16,154	276 3 171 4 98	261 14 183 7 57	11 8 1 2	2 1	1	16 1 3 4 8
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	4,831 26 49 30 1,509 508 1,920 125 664	4,905 21 40 13 1,202 525 2,349 103 652	105 2 3 61 14 10 11	100 4 4 34 15 17 18 5 3	18 3 1 7 1 2 4	28 	1 1 1	6 1 1 1 1 1 2
PACIFIC Nash. Oreg. Calif. Alaska Hawaii	9,852 1,153 345 8,032 160 162	11,090 1,056 469 9,187 155 223	81 13 17 49	74 9 11 54	34 11 N 23	40 9 N 30	51 3 3 45 N	65 2 6 57 N
Guam P.R. VI. Amer, Samoa C.N.M.I.	275	31 172 U U	i	0		Û	N	

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States, weeks ending July 8, 2000, and July 10, 1999 (27th Week)

						Salmon	ellosis*	
Maleria   Rabies, Ar   Cum.   Cum.			NET.			ILIS		
Reporting Area				Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999
NITED STATES				3,040	14,107	15,741	9,816	14,598
Aaine I.H. It. Aass, I.I.	4 1 2 6 4	2 1 11 2	73 8 34 117 21	427 79 26 60 94 51 117	932 68 64 57 530 40 173	936 61 49 37 535 52 202	871 38 62 58 488 49 176	965 50 60 36 529 72 218
IID. ATLANTIC Ipstate N.Y. I.Y. City I.J. a.	86 30 29 9	170 34 82 34 20	492 340 U 80 72	563 392 U 102 69	1,790 523 390 418 459	2,152 502 627 481 542	1,748 502 560 307 379	2,022 525 651 469 377
.N. CENTRAL Ohio nd. II. Aich. Vis.	54 12 3 19 15 5	81 9 8 35 20 9	30 9 1 20	49 12 2 25 10	2,066 555 255 605 421 230	2,422 451 206 802 466 497	1,241 423 233 1 428 156	2,101 451 206 743 462 239
V.N. CENTRAL Winn. owa Wo. V. Dak. S. Dak. Nebr. Cans.	23 8 1 4 2	26 5 7 10	270 48 42 11 74 48	405 54 65 14 84 120 3	980 201 151 347 27 36 64 154	976 221 102 345 15 44 108 141	1,012 282 94 397 39 37 44 119	1,113 334 95 402 30 68 83 101
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fla.	140 3 45 8 28 1 11 1 4 39	150 1 47 10 31 1 10 2 13 36	1,174 20 227 284 63 287 70 157 66	1,089 30 232 271 65 221 79 101 90	2,834 48 385 29 389 72 386 272 450 803	3,082 56 360 46 534 59 469 188 486 884	1,791 51 339 U 302 67 292 180 514	2,608 65 377 U 486 63 517 165 670 265
E.S. CENTRAL (y. Tenn. Ala. Miss.	19 5 5 8	12 2 5 4	94 14 48 32	151 24 55 72	740 165 190 220 165	826 185 211 236 194	432 111 194 111	598 134 227 204 33
W.S. CENTRAL Ark. La. Okla. Tex.	7 1 2 4	13 2 9 2	36	62	1,090 218 107 155 610	1,414 189 291 170 764	1,264 105 177 104 878	1,169 76 254 125 714
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	23 1 1 1 1 3 3 4	22 4 1 1 9 2 2 2 1	113 34 1 26 10 39 2	103 37 29 1 4 31	1,306 58 69 24 392 108 354 172 129	1,425 28 41 20 409 216 402 218 91	908 14 363 83 267 181	1,291 1 46 22 400 166 365 243 49
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	122 12 22 86	127 10 13 96	130 2 109 19	191 1 183 7	2,369 213 163 1,865 29 99	2,508 224 244 1,812 23 205	549 237 202 18 92	2,731 427 272 1,854 13 165
Guam P.R. V.I. Amer, Samoa C.N.M.I.	:	000	32	50 U	109	20 249 U U	00000	0000

N: Not notifiable. U: Unavailable. .: No reported cases.

\* Individual cases can be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health Laboratory Information System (PHLIS).

TABLE II. (Cont'd) Provisional cases of selected notifiable diseases, United States,

	WCCK3 C	Shigeli		o, and so		99 (27th W	GCK/	
	NÉT:			ILIS		Secondary)	Tuber	rculosis
Reporting Area	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999'
INITED STATES	8,468	6,952	3,994	3,957	2,964	3,401	5,309	7,679
HEW ENGLAND Maine H.H. ft. Mass. L.I. Conn.	169 6 3 1 120 12 27	175 3 7 4 116 14 31	131 6 86 12 27	150 6 3 98 9	41 1 1 31 3 5	31 1 2 19 1 8	192 2 4 2 116 23 46	204 11 4 116 20 53
MID. ATLANTIC Jpstate N.Y. V.Y. City V.J. Pa.	1,015 425 387 120 83	480 120 157 124 79	624 146 326 76 76	282 34 120 102 26	121 7 40 24 50	151 13 66 32 40	1,172 131 637 280 124	1,198 141 639 258 160
E.N. CENTRAL Dhio nd. II. Mich. Wis.	1,861 143 760 431 402 125	1,318 260 69 486 169 334	527 95 51 2 346 33	632 63 29 386 122 32	584 39 220 168 136 21	609 50 209 220 113 17	604 132 40 305 82 46	771 108 59 390 163 51
W.N. CENTRAL Minn. Iowa Mo. N. Dak. S. Dak. Nebr. Kans.	895 189 241 351 4 2 25 83	565 82 9 410 2 8 31 23	603 201 131 221 3 1 9	391 103 13 226 2 5 23 19	37 3 10 19 2 3	80 7 7 52 4 10	236 79 23 94 2 9 10	260 100 26 97 2 3 12 20
S. ATLANTIC Del. Md. D.C. Va. W. Va. N.C. S.C. Ga. Fia.	1,220 8 67 16 201 3 64 63 122 676	1,119 8 62 31 42 5 115 62 108 686	322 6 23 U 133 3 26 46 36	297 3 21 U 23 3 57 36 42 113	1,005 5 143 31 69 1 299 97 178 182	1,113 4 227 26 89 2 250 140 210 165	1,177 134 7 108 18 162 54 214 480	1,608 20 134 27 121 23 212 169 317 585
E.S. CENTRAL Ky. Tenn. Ala. Miss.	428 107 210 23 88	693 130 446 63 54	258 44 200 11 3	452 93 323 35	458 51 286 62 59	601 52 331 131 87	381 58 171 152	490 98 153 151 88
W.S. CENTRAL Ark. La. Okla. Tex.	949 108 71 64 706	1,235 51 104 325 755	973 24 72 16 861	509 21 57 99 332	403 49 100 73 181	515 37 129 110 239	165 101 1 63	1,077 82 U 66 929
MOUNTAIN Mont. Idaho Wyo. Colo. N. Mex. Ariz. Utah Nev.	483 4 31 1 80 47 210 35 75	354 6 5 2 54 45 191 27 24	202 2 37 22 105 36	238 6 1 42 31 120 27	107 1 1 2 15 85	116 1 1 6 102 2 4	233 6 5 1 30 29 102 22 38	229 5 1 U 30 114 25 54
PACIFIC Wash. Oreg. Calif. Alaska Hawaii	1,448 315 95 1,006 7 25	1,013 53 37 900	354 279 55 3 17	1,006 56 32 897	208 35 4 168	185 39 3 141 1	1,149 113 8 909 51 68	1,842 86 57 1,584 31 84
Guam P.R. V.I. Amer. Samoa C.N.M.I.	1	7 54 U U	0000	00000	65	1 85 U U	•	103 U U

N: Not notifiable. U: Unavailable. : No reported cases.
"Individual cases can be reported through both the National Electronic Telecommunications System for Surveillance (NETSS) and the Public Health Laboratory Information System (PHLIS).
"Cumulative reports of provisional tuberculosis cases for 1999 are unavailable ("U") for some areas using the Tuberculosis Information System (TIMS).

TABLE III. Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending July 8, 2000, and July 10, 1999 (27th Week)

	H. influ	ienzae,			10, 1999 iral), By Ty				Meas	es (Ruber	ola)	
		sive	A		В		Indiger	nous	Impo		Tota	1
Reporting Area	Cum. 2000'	Cum. 1999	Cum. 2000	Cum. 1999	Cum. 2000	Cum. 1999	2000	Cum. 2000	2000	Cum. 2000	Cum. 2000	Cum 1999
INITED STATES	642	633	5,575	9,736	3,349	3,482	3	29	12000	10	39	59
EW ENGLAND	47	46	140	119	35	76				3	3	9
Aaine	1	5	10	4	5				-	-		
I.H.	9	7	16	7	10	8	-			*		1
ft.	3	4	4	1	4	1		100		3	3	
Aass. R.I.	23	19	56 7	47	7 9	27 17		*		*	-	6
onn.	10	10	47	51	3	23						2
					007							
AID. ATLANTIC Jpstate N.Y.	101 50	114	335 106	621 132	337 64	481 108	3	6	-	1	7	5
V.Y. City	23	35	150	165	194	145	3	0		-	0	2
V.J.	21	29	79	80	79	73						
Pa.	7	3	-	244		155		+		1	1	
N. CENTRAL	84	100	682	1,635	364	346		6			6	2
Ohio	35	37	146	376	65	49		2			2	
nd.	11	14	33	59	26	27					8	1
(1,	33	41	238	352	61			3		-	3	
Mich.	5	8	252	802	211	247		1		-	1	1
Nis.			13	46	1	23	-	~	1.5			
W.N. CENTRAL	36	26	589	394	492	144		2	+	1	3	
Minn.	16	13	129	35	19	22			-	1	1	
owa	8	1	49	74	21	23		1	*		1	
Ио. N. Dak.	1	3	284	233	407	84	Ü		1.0	-	*	
S. Dak.		2	2	8	2	1	U		U		-	
Vebr.	4	3	18	33	18	11	U		U			
Cans.	7	4	107	10	25	3	-	1		-	1	
S. ATLANTIC	177	138	698	923	636	542		1			1	4
Del.	111	130	030	2	030	342					1	-
Ad.	46	34	84	173	68	97				*		
D.C.	-	4	11	35	16	12						
Va.	28	12	76	82	78	53		*				5
W. Va.	5	4	44	19	6	14		~	-	*		
N.C. S.C.	15	22	92 30	64 21	139	125 38	1				*	
Sa.	50	41	111	268	98	63						
Fla.	25	19	250	259	226	140		1			1	
S. CENTRAL	29	42	234		244	240						
Cy.	11	6	27	233	50	246		-			~	-
lenn.	13	21	90	97	109	117				*		
Ala.	4	13	31	36	27	52	-			+	-	
Miss.	1	2	86	56	58	58		-			*	
W.S. CENTRAL	36	42	925	2,898	353	590		1			1	
Ark.		1	89	25	56	43		1			1	
.a.	7	11	28	97	50	113	-		-		-	
Okla.	27	27	153	303	71	73					4.1	
Tex.	2	3	655	2,473	177	361	-6	00		- 20	-	4
MOUNTAIN	69	57	470	734	250	325	*	11		1	12	
Mont.		1	2	12	3	16						
daho Wyo.	3	1	17	29	5	18	*	-	*	~		
rvyo. Colo.	11	9	100	142	51	47		1		1	2	
N. Mex.	14	13	41	30	64	104					2	
Ariz.	33	28	235	417	89	80						
Jtah	6	2	34	28	14	20		3			3	
Nev.	1	2	33	72	22	33		7		*	7	
PACIFIC	63	69	1,502	2,179	638	732		2		4	6	3
Nash.	3	2	145	168	41	34		-		-	-	-
Oreg.	18	24	116	147	50	61			*			1
Calif.	24	36	1,233	1,847	537	616		1		2	3	3
Alaska Hawaii	2 16	5 2	8	13	5	13		1	-	2	1	
	10	2			9	8				2	2	
Guam	-			2	-	2	U		U	-		
P.R. V.L	1	2 U	55	173	54	125	Ü		U	*	-	
Amer. Samoa		Ü		U		U	Ü		Ü			1
C.N.M.I.		ŭ		ŭ		U	Ü		Ü	-	-	1

N: Not notifiable. U: Unavailable. -: No reported cases.
\*For imported measles, cases include only those resulting from importation from other countries.
'Of 130 cases among children aged <5 years, serotype was reported for 59 and of those, 16 were type b.

TABLE III. (Cont'd) Provisional cases of selected notifiable diseases preventable by vaccination, United States, weeks ending July 8, 2000, and July 10, 1999 (27th Week)

	Mening Disc	ococcal ease		Mumps			Pertussis			Rubella	
Reporting Area	Cum. 2000	Cum. 1999	2000	Cum. 2000	Cum. 1999	2000	Cum. 2000	Cum. 1999	2000	Cum. 2000	Cum 1999
JNITED STATES	1,229	1,418	4	198	217	44	2,485	3,010	19	76	160
IEW ENGLAND	77	69		2	5	4	636	331		6	7
Maine	6	5		~	-		14	-		-	
I.H. /t.	9 2	9		*	1	4	62 142	53 25		2	
Aass.	47	41			4		380	236		3	7
R.I.	5	2	*	1	*		9	8			
Conn.	8	8	*	1			29	9	. "	1	
MID. ATLANTIC	119 36	144 39	*	9	29 5	6	189	597 499	-	2 2	23 15
Jpstate N.Y. N.Y. City	25	42		0	7	6	115	19		2	2
V.J.	26	31	-	-	1			15			3
Pa.	32	32	*	3	16		74	64			3
.N. CENTRAL	221 51	253 94		23 7	27	1	279	253 114		1	2
Ohio nd.	34	32		,	3		167 27	14	-		1
II.	53	65		5	7		23	54	*	1	1
Mich. Wis.	64 19	35 27	*	11	8	1	31 31	25 46			
							135				
W.N. CENTRAL Minn.	104	139 29	-	12	9	1	66	109	-	1	83
owa	19	26	*	5	4	*	24	23			25
Mo. N. Dak.	61	51	ű	1	1	ú	23	27	ú		2
S. Dak.	5	8	0	-	- 2		3	4		-	
Nebr.	5	8	U	2	3	U	3	3	U	1	56
Cans.	5			4		1	15	19		1	-
S. ATLANTIC Del.	201	219	1	32	36	15	208	154	19	51	20
Md.	19	35		7	4	2	47	47		-	1
D.C. Va.	32	1 26		5	2		21	13		*	
W. Va.	8	4	-	-	-	1	1	1		-	
N.C. S.C.	30	27	1	5	8	2	51	42	19	42	19
Ga.	15 33	30 43		10 2	3		19 20	8 16		7	
Fla.	64	49		3	9	9	43	27		2	
E.S. CENTRAL	88	108		6	7	2	45	54		4	1
Ky. Tenn.	18	19 41	*	2		1	19 14	12 27		1	
Ala.	25	29	-	2	5	1	11	13	2	3	
Miss.	7	19		2	2		1	2			
W.S. CENTRAL	87	144	1.00	20	29	2	117	85		4	
Ark. La.	9 27	26 52	-	3	6		10	8	*		
Okla.	21	21			1		6	8			
Tex.	30	45		16	22	2	98	64		4	
MOUNTAIN	66	89	*	14	10	8	407	364		2	15
Mont. Idaho	6	2 8		1	1	1	9	160			
Wyo.		3		1		-	3	2			
Colo. N. Mex.	24	22 11		1	3	5	225 74	131 36		1	
Ariz.	18	29		3	N	1	41	60		1	10
Utah	7	9	*	4	3	-	9	31			
Nev.	3	5			3		6	2			
PACIFIC Wash.	266 32	253 38	3	80	66	5	469 181	1,063 505	*	5	
Oreg.	36	47	N	N	N	2	56	21			
Calif.	187	158	3	67	56		215	510		5	
Alaska Hawaii	5	6		3	7		12	3 24			
Guam		1	U		1	U		1	U		
P.R.	5	9	U			U		13	U		
V.I. Amer. Samoa	*	U	U		U	U	*	U	U		1
C.N.M.I.	-	Ü	Ü		Ü	Ü		ŭ	Ü		-

II: Not notifiable.

U: Unavailable.

TABLE IV. Deaths in 122 U.S. cities,\* week ending

	All Causes, By Age (Years)									All Causes, By Age (Years)					
	All	:65	45-64	25-44	1-24	<1	Total	Reporting Area	All Ages	≥65	45-64	25-44	1-24	<1	P&I Tota
NEW ENGLAND Joston, Mass, Zridgeport, Conn. Joseph Green, Mass, Joseph Green, Mass, Joseph Green, Mass, New Bedford, Mass, New Haven, Conn. Providence, R.I. Somerville, Mass, Springfield, Mass,	37 U 6 25	358 77 40 10 17 35 29 12 18 24 U 4 22 29	2 6 14 2 9 U	29 13 2 4 1 1 1 U	9 3 1 2 1	6 2 2 1	42 9 1 1 1 8 5 1 1 1 1 0 7	S. ATLANTIC Atlanta, Ga. Baltimore, Md. Charlotte, N.C. Jacksonville, Fla. Morfolk, Va. Richmond, Va. Savannah, Ga. St. Petersburg, F Tampa, Fla. Washington, D.C. Wilmington, Del	17 8 13 10 4 4 14 15 16 17 18 18 18 18 18 18 18 18 18 18 18 18 18	U U 106 10 46 15 91 14 55 14 55 16 37 25 37 25 38 88	U U 38 38 16 16 31 32 33 5 9 10 4 4 20 28 7 27	72 U 22 12 8 13 1 5 1	21 U 2 1 1 5 2 1 2 3 4	20 U 3 5 4 2 1 1	560 U 133 44 99 88
Waterbury, Conn. Worcester, Mass. MID. ATLANTIC Allbany, N.Y. Allentown, Pa. Buffalo, N.Y. Zamden, N.J. Litzabeth, N.J. Frie, Pa.§	34 57 2,079 42 U 100 28 5 36	41 1,396 34 U 69 15 4 31	12 424 3 U 20 4 1	166 2 U 6 5	51 2 U 3 2	40 1 U 2 2	7 110 3 U 11	E.S. CENTRAL Birmingham, Ala Chattanooga, Te Knoxville, Tenn. Lexington, Ky. Memphis, Tenn. Mobile, Ala. Montgomery, Al Nashville, Tenn.	nn. 2	14 91 38 55 77 55 30 1	31 8 2 8 2 17 3 14 4 53 1 15 3 4	70 7 5 4 4 16 21 1	22 7 2 2 1 4 4	22 1 1 2 1 9 4	56 20
Jersey City, N.J. New York City, N.Y. Newark, N.J. Paresron, N.J. Philadelphia, Pa. Pittsburgh, Pa.S. Reading, Pa. Rochester, N.Y. Scranton, Pa.S. Syracuse, N.Y. Trenton, N.J. Ulica, N.Y. Yonkers, N.Y.	41 1,104 52 10 299 37 39 127 22 33 58 29 18 U	20 725 24 6 197 25 34 95 16 28 45	246 16 2 62 7 2 2 2 2 4 4 7 9 3	6 91 5 2 26 2 2 6 2 1 3 5	3 20 4 13 1	2 21 3 1 3 2	45 1 23 1 4 10 1 1 3 2	W.S. CENTRAL Austin, Tex. Baton Rouge, La Corpus Christi, Dallas, Tex. El Paso, Tex. Ft. Worth, Tex. Houston, Tex. Little Rock, Ark. New Orleans, La San Antonio, Te Shreveport, La. Tulsa, Okla.	Tex. 10	68 4 26 1 47 3 54 9 51 3 98 6 13 17 39 2 U 1 83 12	6 12 8 6 2 8 2 41 3 12 7 23 9 69 4 10 U U 0 38 3 11	6 19 5 6 44 3 U 15 6	47 3 1 7 2 16 1 1 0 7 6 4	22 1 1 5 1 5 1 U	10:
E.N. CENTRAL Akron, Ohio Canton, Ohio Chicago, Ill. Cincinnati, Ohio Cleveland, Ohio Columbus, Ohio Dayton, Ohio Detroit, Mich. Evansville, Ind. Fort Wayne, Ind.	1,415 42 38 393 U 111 134 84 150 37 55	923 31 25 254 66 83 57 72 26 31	5 6 4 84 5 26 3 28 7 19 2 42 6 7 3 13	36 U 11 17 6 17 2	35 2 9 U 3 4 1 7	40 2 10 U 5 2 1 12 1	97 2 3 33 U 5 6 5 7	MOUNTAIN Albuquerque, N Boise, Idaho Colo. Springs, C Denver, Colo. Las Vegas, Nev. Ogden, Utah Phoenix, Ariz. Pueblo, Colo. Salt Lake City, U Tucson, Ariz.	.M. lolo. 1	65 4 84 4 78 11 29 2 51 9 15 1 84 5	8 24 5 5 7 12 2 17	7 1 3 14 13 3 16 2 7	7	20 1 9 2 6 1 1	1
Gary, Ind. Grand Rapids, Mich Indianapolis, Ind. Lansing, Mich. Milwaukee, Wis. Peoria, III. Rockford, III. South Bend, Ind. Toledo, Ohio Youngstown, Ohio	U 36 79 26 43 31 102	10 27 27 55 14 32 77	7 4 J U 2 9 7 19 4 7 5 2 5 5 7 18	2 U 3 2 3 3 1 1 6	3 U 1 1 1	1 1 2 1 1 U	5 U 2 4 2 6 1	PACIFIC Berkeley, Calif. Fresno, Calif. Glendale, Calif. Honolulu, Hawa Long Beach, Cal Los Angeles, Ca Pasadena, Calif. Portland, Oreg. Sacramento, Ca	lif. lif. 1,0	14 75 47 58 67 04 68 27	11 2 12 15 10 8 18 7 18 9 12 211 21 3	5 7 2 3 71 71 71 3 1	2 1 3 26 1	1 4 14 1	15
W.N. CENTRAL Des Moines, lowa Duluth, Minn. Kansas City, Kans. Kansas City, Mo. Lincoln, Nebr. Minneapolis, Minn Omaha, Nebr.	784 U 13 90 96 21 1. 130 63	9	U U U 1 B 19 0 20 6 4 0 22	U 2 8 11 11 1 9	25 U -4 5	-	4 4	San Diego, Calit San Francisco, C San Jose, Calif Santa Cruz, Cali Seattle, Wash. Spokane, Wash Tacoma, Wash.	f. 1 Calif. 2 f. 1	32 U 00 14 21 U 56 4	77 32 U U 12 37 15 5 U U 10 7 33 20	2 6 3 U 12 5 1 3 U 7 3	2 U 5 U 2 5	5 U 4 U 4 1	
St. Louis, Mo. St. Paul, Minn. Wichita, Kans.	96 45 260	3	1 13	8	2 1 9	2	6	TOTAL	10,5	034 6,96	59 2,134	833	290	227	7

U: Unavailable. ∴No reported cases.

\*Mortality data in this table are voluntarily reported from 122 cities in the United States, most of which have populations of ≥100,000. A death is reported by the place of its occurrence and by the week that the death certificate was filed. Fetal deaths are not included. 
Pneumonia and influenza.

\*Because of changes in reporting methods in this Pennsylvania city, these numbers are partial counts for the current week. Complete counts will be available in 4 to € weeks.

\*Total includes unknown ages.

Since July 1999, substantial progress has been made in removing thimerosal from vaccines. As of March 2000, all U.S. children had access to hepatitis B vaccines that do not contain thimerosal as a preservative. Beginning July 2000, only single-dose thimerosal-free *Haemophilus influenzae* type b vaccine will be produced in the United States; previously manufactured multidose vials containing thimerosal still may be in distribution. One diphtheria and tetanus toxoids and acellular pertussis vaccine (DTaP) that does not contain thimerosal is available, and it is projected that additional DTaP vaccines without thimerosal as a preservative will become available in early 2001. On the basis of this progress, the most likely maximum amount of ethylmercury that an infant may be exposed to from the routine vaccination schedule has been reduced by 60%, from 187.5 µg to 75 µg. Measles-mumps-rubella, varicella, inactivated polio, and pneumococcal conjugate vaccines have never contained thimerosal.

Research on the potential health effects of exposure to thimerosal is continuing, and findings will be monitored closely by PHS to determine whether any changes in policy are needed. AAFP, AAP, and PHS, in consultation with the ACIP, reaffirm the goal set in July 1999 to remove or greatly reduce thimerosal from vaccines as soon as possible. On the basis of information from the Food and Drug Administration and manufacturers, PHS projects that the United States will complete its transition to a secure routine pediatric vaccine supply free of thimerosal as a preservative by the first quarter of 2001.

The vaccination of children in much of the world will continue to require the use of multidose vials because of cost, production, and storage capacity. Multidose vials require a preservative to prevent microbial contamination after the vial is opened. For multidose vials, manufacturers are encouraged to seek alternatives to thimerosal.

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